

**REMARKS****Claim Status**

Claims 1-17, 19-33, 37 and 38 are pending in this application and stand rejected. Claims 13, 14, 17, 33 and 38 are canceled herein without prejudice. Claims 1, 3, 16, 31 and 37 are amended herein, and new claims 44 and 45 have been introduced. Support for the claim amendments and new claims is provided by the specification at, e.g., page 7, lines 13-18; page 10, lines 15-21; page 31, lines 18-25; page 33, lines 3-7; page 34, lines 13-17; and throughout the Examples. Accordingly, no new matter is added by way of these amendments.

The instant claim amendments are marked relative to the last official claim listing entered and examined by the Office, submitted with the response filed on March 22, 2010. This listing of claims will replace all prior versions and listings of claims in this application.

Applicants respectfully request entry of the claim amendments and new claims and reconsideration in view of the following remarks.

**Rejections under 35 U.S.C. § 103**

Claims 1-17, 19-33, 37 and 38 stand rejected under 35 U.S.C. § 103(a) as allegedly unpatentable over WO 98/34644, for reasons of record, in view of Chen (U.S. 6,602,274). Briefly, WO 98/34644 is said to disclose that low dose PDT can be used to reduce inflammation in injured or pre-injured tissues, and Chen is said to disclose that PDT causes damage to normal tissues beyond the treated area. Applicants traverse the rejections for reasons of record, as well as at least the following reasons.

Independent claim 1 is amended herein to clarify that the area treated with low dose PDT in step b) of the method encompasses (i.e., includes) the target tissue that is treated with normal dose PDT in step a). Independent claim 31 is similarly amended to clarify that the area treated with low dose PDT in step d) encompasses (i.e., includes) the treatment area that is treated with normal

dose PDT. Therefore, the claims cannot be mistakenly construed to mean that the low dose PDT is delivered only to a “doughnut” shaped area around the target tissue.

Thus, it will be understood that the “target tissue” in claim 1 and the “treatment area” in claim 31 are exposed to a total light dosage that is the sum of light administered in both the normal dose and the low dose PDT treatment steps. This total dosage is necessarily significantly higher than the dosage of the low dose PDT treatment alone. Both claims 1 and 31, as amended, clearly require that the target tissue (claim 1) or treatment area (claim 31) is exposed to a total light dose that is the sum of the normal dose PDT and the low dose PDT, which is necessarily significantly higher than that of low dose PDT alone.

WO 98/34644 describes appropriate irradiation conditions for low dose PDT as: irradiation at a dose of less than 15 J/cm<sup>2</sup> applied between 0-3 hours after administration of the photosensitizer, when significant amounts of the photosensitizer would still be expected to be present (page 36, lines 24-25). Irradiation at light doses up to 100 J/cm<sup>2</sup> is described when irradiation is applied later than 6 hours after photosensitizer administration, at which time most of the photosensitizer would be expected to have been cleared (page 36, lines 26-27). Particularly preferred conditions for use during surgery are said to include exposing the treated tissue to about 7-12 J/cm<sup>2</sup> of light (page 37, line 20).

In addition, WO 98/34644 at page 40, lines 3-9 states that: “The data indicated that a certain level of PDT was required, but that higher doses were generally less effective than lower ones. The combination of the short incubation time with BPD and the low light dosage of 12 J/cm<sup>2</sup> was not expected to cause much damage to treated cells. Nevertheless, the treatment had a definite pharmacological action. Bleb survival was associated with the lack of inflammation, as indicated by avascularity and a pale-colored bleb.”

The data presented in Table 1A and 1B on page 39 of WO 98/34644 shows that irradiation above 12 J/cm<sup>2</sup> resulted in vascularity, and poor bleb survival, indicators of inflammation (see, e.g., Table 1A and B, rabbit nos. 5 and 6, treated with 18 and 24 J/cm<sup>2</sup>, respectively).

Applicants respectfully submit that these statements and data would have suggested to one of ordinary skill that there is a maximum PDT dose that will provide the anti-inflammatory effect and that at higher irradiation doses, inflammation occurs. Taken as a whole, WO 98/34644 suggests that irradiation at a dose of less than  $15 \text{ J/cm}^2$  would be preferred, and that irradiation at higher light doses did not provide anti-inflammatory efficacy.

As amended, both claims 1 and 31 require that the target tissue (claim 1) or treatment area (claim 31) is exposed to a total light dose that is higher than the low dose PDT indicated by WO 98/34644. In view of the disclosure of WO 98/34644, one of ordinary skill in the art would not have reasonably believed that irradiation at a total light dose greater than the low dose PDT reported by WO 98/34644 would have provided the anti-inflammatory effects described by the cited reference at lower light doses.

Accordingly, Applicants respectfully submit that one of skill in the art would not have had a reasonable expectation of success that the instantly claimed methods would be effective to reduce or prevent inflammation in a treated subject. To the contrary, Applicants submit that a skilled person would be surprised at the discovery of reduced inflammation by using a total combined PDT dose that exceeds the low dose PDT reported in WO 98/34644. Applicants further submit that in view of the lack of a reasonable expectation of success and the unexpected nature of the invention, no basis has been provided to explain why one of skill in the art would have made the modifications to the cited art necessary to achieve the instant invention.

In view of the foregoing remarks, Applicants respectfully request that the rejections under 35 U.S.C. § 103(a) be withdrawn.

**CONCLUSION**

In view of the above, each of the presently pending claims in this application is believed to be in immediate condition for allowance. Accordingly, the Examiner is respectfully requested to withdraw the outstanding rejection of the claims and to pass this application to issue. If it is determined that a telephone conference would expedite the prosecution of this application, the Examiner is invited to telephone the undersigned at the number given below.

In the event the U.S. Patent and Trademark office determines that an extension and/or other relief is required, applicant petitions for any required relief including extensions of time and authorizes the Commissioner to charge the cost of such petitions and/or other fees due in connection with the filing of this document to Deposit Account No. 03-1952 referencing docket no. 273012011800. However, the Commissioner is not authorized to charge the cost of the issue fee to the Deposit Account.

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